

Gastrointestinal Stromal Tumors: Clinical Symptoms, Location, Metastasis Formation, and Associated Malignancies in a Single Center Retrospective Study

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Keywords

Gastrointestinal stromal tumor · Gastrointestinal oncology · Gastrointestinal symptoms · Gastrointestinal tract · Metastasis · Recurrence

Abstract

Background and Aims: Gastrointestinal stromal tumors (GISTs) are rare malignancies but the most common mesenchymal tumors of the digestive tract. Recent advances in diagnostic imaging and an increasing incidence will confront us more frequently with stromal tumors. This single center study aimed to characterize GIST patients in terms of tumor location, clinical presentation, metastasis formation, as well as associated secondary malignancies. **Methods:** In a retrospective study, 104 patients with a histologically confirmed diagnosis of GIST, collected between 1993 and 2011, were characterized for several clinical features. **Results:** The most common GIST location was the stomach (67.6%) followed by the small intestine (16.2%). Gastrointestinal bleeding (55.8%) and abdominal pain (38.5%) were the most frequently reported symptoms whereas about one-third of patients remained clinically asymptomatic (31.6%); 14.4% of patients had either synchronous or metachronous metastases and

there was a significant prevalence also in the low risk group. The proportion of secondary malignant associated neoplasms was 31% in our GIST cohort, among which gastrointestinal, genitourinary tumors, and breast cancer were the most prevalent. **Conclusion:** There was a considerable risk for metastasis formation and the development of secondary neoplasias that should encourage discussion about the appropriate surveillance strategy after surgery for GIST.

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Introduction

With only 0.1–3% of all gastrointestinal neoplasms, gastrointestinal stromal tumors (GISTs) are a rare tumor entity. Nonetheless they represent the most common mesenchymal tumors of the digestive tract and are even found in extraintestinal locations (eGIST) [1, 2]. Arising from the interstitial cells of Cajal or their precursors, they show a set of characteristic features including a spindle cell, epithelioid or, rarely, a pleomorphic morphology.

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Mutations in the genes for cKIT (CD117; 85% of all GIST) and platelet-derived growth factor receptor alpha (5–7%) were ascribed a central role in the pathogenesis by permanent tyrosine kinase activation [3–5].

The annual incidence of GISTs ranges from 0.7 to 2 per 100,000 and increased during the last years [6, 7]. Data from the Surveillance, Epidemiology, and End Results registry from the U.S. National Cancer Institute found an age-adjusted yearly incidence of 6.8 per million [8, 9].

Several risk classifications systems exist for stratification of stromal tumors such as those by Fletcher et al. [10], the modified National Institutes of Health (NIH) classification by Joensuu et al. [11], or the TNM-classification [12]. The ESMO guidelines favor the use of a classification by the Armed Forces Institute of Pathology (AFIP) [6] that considers tumor size, location, and their mitotic rate as prognostic factors [13]. Modified NIH criteria encompass a fourth factor, which is tumor rupture [14].

In early stages, stromal tumors often remain asymptomatic and are detected accidentally. Clinical symptoms vary and include gastrointestinal bleeding (51%), a palpable mass (36%), and abdominal pain (33%) whereby the stomach is the most common site of disease manifestation followed by the small intestine [15]. Usually diagnosis is achieved using a combination of endoscopy, laparoscopy, and computed tomography [6]. Treatment options are manifold depending on the size and location of the tumor, localized or metastasized disease status, age, and comorbidities of the patient and include surgical and medical procedures. Standard treatment comprises surgical excision unless oncologic resection is feasible and no major comorbidity exists. In case of a high estimated risk for GIST recurrence after surgery, adjuvant therapy with tyrosine kinase inhibitors (imatinib) is initiated [16]. Mutational analysis is critical as some genotypes limit the success of adjuvant therapy such as in platelet-derived growth factor receptor alpha D842V-mutated GISTs. In locally advanced and metastasized stages, imatinib is the standard treatment and dose modification may be required according to the genotype of the KIT gene. In case of tumor progression or intolerance, sunitinib and regorafenib are considered as second- and third-line treatment options [6, 17, 18].

As known for other malignant tumors, GISTs harbor a risk of relapse and metastasis formation [19]. Secondary tumors, mainly gastrointestinal malignancies, have been reported in association with stromal tumors, which further worsen prognosis [20, 21].

Here, we present data on GIST patients that were diagnosed in a single center in Western Pomerania, a rural region of Germany, in a time period from 1993 to

2011. We focused on clinical symptoms and their presence depending on tumor location. Secondly, we investigated the occurrence of metastases and the prevalence of associated secondary tumors.

Patients and Methods

Selection of Patients

After ethical approval from the local institutional review board committee, we collected clinicopathological and follow-up data of patients with the diagnosis of a GIST at University Medicine Greifswald in a retrospective monocentric analysis between 1993 and 2011. Patients were identified in the hospital information system using the ICD-10 codes C15 to C18 and C48, and via cross-checking of histology reports of the Institute of Pathology. A total of 104 patients were identified with residence in Western Pomerania (urban district of Greifswald, Vorpommern-Greifswald, Vorpommern-Rügen, and Mecklenburg-Lake counties). The diagnosis of GIST was histologically confirmed for all patients and was done in all cases by the Institute of Pathology of University Medicine Greifswald. From 1993 to 1998, diagnosis of GIST was established retrospectively because this tumor was defined as an independent entity in 1998 [22]. Tumors were immune-stained for CD117, CD34, and smooth muscle antigen. Specimens from 1993 to 1998 were retrospectively stained with CD117 and CD34. The number of residents in Western Pomerania (the same catchment area) was obtained from the statistical office of Mecklenburg-Western Pomerania based on the census of 2013 [23]. The number of inhabitants was 727,270 on January 1, 2013.

Acquisition of Clinical Data and Statistical Analysis

After identification of patients, they were contacted by mail and informed about the intention and type of study. A written informed consent was obtained from all still living individuals. In those patients who already died, consent was obtained by proxy and data fully anonymized. Patient data included gender, age at diagnosis of GIST, clinical symptoms, size, localization and genotype of the tumor, diagnostic examinations performed, therapy, and follow-up data. Histological subtypes of GIST and proliferation index were assessed, too. Risk of tumor recurrence was stratified according to the AFIP-Miettinen classification system, the Fletcher consensus criteria, and the modified NIH classification by Joensuu, whenever possible. Results were reported as percentages, mean \pm SD, and median and range, when applicable. A 2-tailed p value of <0.05 was considered to be significant. Data were collected using Microsoft® Access 2013 (Redmond, WA, USA) software and have been analyzed by the translational research platform tranSMART in version 16.2 [24].

Results

Characterization of the Cohort and Pathology of the Tumor

The study included 104 patients showing an almost equal distribution of female (48.1%) and male (51.9%) patients. Mean age of first diagnosis of GIST was 66.86 years

Table 1. Characterization of the study group

Quality	Results
Age and gender	
Patients with diagnosis of GIST	104
Age at diagnosis, years, mean \pm SD (range)	66.86 \pm 11.85 (30–88)
Males, <i>n</i> (%)	54 (51.9)
Tumor size and histology	
Tumor diameter available	94
Tumor size, cm, mean \pm SD (range)	4.96 \pm 3.85 (0.2–20)
Histology available	93
Spindle cell, <i>n</i> (%)	72 (77.4)
Epithelioid cell, <i>n</i> (%)	12 (12.9)
Mixed type, <i>n</i> (%)	9 (9.7)
Clinical symptoms	
Data of clinical symptoms available	76 (73.1% of all 104 patients)
Clinical symptoms present, <i>n</i> (%)	52 (68.4)
Clinically asymptomatic, <i>n</i> (%)	24 (31.6)
Metastases	
Patients with metastases	15
Synchronous metastases, <i>n</i> (%)	5 (33.3)
Metachronous metastases, <i>n</i> (%)	10 (66.6)
Mean time of recurrence, years	3.6
Secondary neoplasia, <i>n</i> (%)	
Secondary neoplasia in addition to GIST	44 (42.3)
Malignant neoplasia	32 (72.7)
Benign neoplasia	14 (31.8)
Malignant and benign neoplasia	3 (6.8)

(SD 11.85 years) and median age was 70 years. We calculated the approximate annual incidence of GISTs between 1999 and 2010 based on the population in Western-Pomerania with a mean of 1.1 per 100,000 and a variance of 0.4–2.6 per 100,000. Mean follow-up for all patients was 30.9 months and was carried out for up to 180 months.

The mean tumor size in our cohort was 4.96 cm (range from 0.2 to 20 cm). Histologically, more than 75% of all tumors showed a spindle cell configuration whereas epithelioid cell and mixed type pattern were found more rarely (Table 1).

Localization of GISTs

Data were available for almost all patients (*n* = 102). One patient had both gastric and an extraintestinal GIST (diaphragm). GISTs were widely distributed along the entire gastrointestinal tract. By far, the most frequent location was the stomach where around two-thirds of all tumors were found. It was found that 16.2% of all GISTs were located in the small intestine whereas only 2.9% of the tumors were found in the colorectum. Esophageal location was seen in 4.8%. Extraintestinal manifestation was observed for 7 tumors corresponding to 6.7% of all GISTs (Table 2).

Table 2. Localization of GIST in the gastrointestinal tract and extraintestinal location

Localization of GIST	Number	Percentage
Stomach	71	67.6
Small intestine	17	16.2
Colorectum	3	2.9
Esophagus	5	4.8
eGIST	7	6.7
No data available	2	1.9
Total	105*	100

* Multiple locations found in 1 patient.

GIST, gastrointestinal stromal tumor; eGIST, extraintestinal eGIST.

Risk Stratification

Based on tumor size, localization, and proliferation rate (mitotic index), tumor stage classification and risk assessment were performed. We used the classification systems of the AFIP [13], NIH [10], as well as the modified NIH criteria [11] and were able to classify 89 patients

Table 3. Recurrence risk according to the classifications of Fletcher et al. [10], Miettinen et al. [13], and Joensuu et al. [11]

Recurrence risk	NIH (Fletcher et al. [10], 2002), n (%)	AFIP (Miettinen et al. [13], 2006), n (%)	Modified NIH (Joensuu et al. [11], 2008), n (%)
Very low	14 (15.7)	17 (19.5)	17 (19.5)
Low	30 (33.7)	43 (49.4)	28 (32.2)
Intermediate	23 (25.8)	6 (6.9)	18 (20.7)
High	22 (24.7)	21 (24.1)	24 (27.6)
Total	89 (100)	87 (100)	87 (100)

by NIH criteria and 87 by modified NIH and AFIP criteria, respectively. When classifying according to the most widely used AFIP system, more than half of patients were grouped to a low (49.4%) or even very low (19.5%) tumor recurrence risk (Table 3). The percentage of patients with a very low recurrence risk was comparable in all 3 classification systems. Low recurrence risk was more frequently seen (49.4%) when applying the AFIP criteria compared to the modified NIH criteria (32.2%) or the initial consensus approach by Fletcher et al. [10] (33.7%). In contrast, only 6.9% of patients had an intermediate tumor recurrence risk according to the AFIP criteria which was remarkably higher (20.7 and 25.8%, respectively) after application of other classification systems. Numbers of patients having a high risk were quite comparable in all 3 classification systems (Table 3).

Clinical Symptoms at First Diagnosis

Data regarding presence of clinical symptoms of GIST patients at hospital admission were available for 76 out of 104 patients (73.1%). Among them 52 patients (68.4%) had symptoms, whereas the remainder was asymptomatic and diagnosis was established during routine examinations or accidentally during a diagnostic workup due to other reasons. We observed a variety of clinical symptoms that were often unspecific and not directly indicating an underlying tumor disease. Gastrointestinal bleeding was the most common reported clinical sign (55.8%), followed by abdominal pain (38.5%), weight loss (13.5%), nausea/vomiting (9.6%), heartburn (9.6%), and dizziness (7.7%; Table 4).

In a next step, clinical symptoms were analyzed in relation to tumor localization in order to clarify whether particular symptoms are associated with a specific tumor site. We only considered patients where both information on GIST localization and clinical symptoms were available (online suppl. Table 1; for all online suppl. material, see www.karger.com/doi/10.1159/000489556).

Table 4. Most frequently reported clinical symptoms of patients with GIST at first diagnosis (multicode allowed)

Symptom	Number of patients, n (%)
Gastrointestinal bleeding	29 (55.8)
Abdominal pain	20 (38.5)
Weight loss	7 (13.5)
Weakness/dizziness	7 (13.5)
Heartburn	5 (9.6)
Nausea/vomiting	5 (9.6)
Loss of appetite	2 (3.8)
Dyspnea	2 (3.8)
Ileus	2 (3.8)
Feeling of fullness	2 (3.8)

Most data on symptoms were available for gastric GISTs since this tumor entity was the most common one. Gastrointestinal bleeding was most frequently reported by patients with GISTs of the stomach (64%), the small intestine (46.7%), and the colorectum (33.3%), although the absolute number of patients with colorectal GISTs was quite low. Regarding tumors of the stomach, abdominal pain was reported by 28% of patients, whereas another 28% of patients did not report any clinical symptoms. Abdominal pain was also associated with tumors of other sites, as in 25% for esophageal, 20% for small intestinal, and 50% for extraintestinal GIST location but tumor prevalence was much lower compared to gastric location.

Metastasis Formation of GISTs

Patients were further analyzed for occurrence of metastases; 15 out of 104 patients were diagnosed with metastases, of which 5 (33.3%) had synchronous and 10 (66.6%) metachronous metastases. Patients with local tumor recurrence after surgery (2 individuals) were also included to the group with metachronous metastases. In all patients with synchronous and metachronous metas-

Table 5. Presence of GIST metastases in relation to risk classifications of Fletcher et al. [10], Miettinen et al. [13], and Joensuu et al. [11]

Recurrence risk	Number of patients, <i>n</i> (%)				
	high risk	intermediate risk	low risk	very low risk	all
NIH (Fletcher et al. [10], 2002)	11 (73.3)	1 (6.7)	2 (13.3)	1 (6.7)	15 (100)
AFIP (Miettinen et al. [13], 2006)	10 (71.4)	0 (0)	3 (21.4)	1 (7.1)	14 (100)
Modified NIH (Joensuu et al. [11], 2008)	10 (71.4)	1 (7.1)	2 (14.3)	1 (7.1)	14 (100)

tases, origin from a GIST was confirmed by histology. The mean time between initial GIST diagnosis and appearance of metastases was 3.6 years (ranging from 1 to 15 years). The most frequent location was the liver (10 of 15 patients, 66.7%), rarer sites included the peritoneum (4 of 15, 26.7%), bones (2 of 15, 13.3%), lymph nodes (2 of 15, 13.3%), and pancreas and the colon (1 of 15, 6.7%).

To determine independent variables for recurrence-free survival, Kaplan-Meier analyses were calculated using the categories tumor location (stomach, esophagus, small intestine, eGIST), size (<5, 5.1–10 and 10.1–15 cm), and mitotic index (0–5, 6–10, >10 mitoses/high power field). Small tumors (<5 cm) and those with a low mitotic index (<5/high power field) were associated with a lower disease-free survival. No clear association was found for tumor location acknowledging a small patient size as a limiting factor (online suppl. Fig. 1).

Next, we compared patients with synchronous or metachronous metastases by using the commonly used risk classifications in order to determine the accuracy of predicting metastatic disease. Again we used the NIH, modified NIH, and AFIP criteria that were available for 15 and 14 patients, respectively (Table 5). As expected, the majority of patients with metastases belonged to the high risk group (71.4 and 73.3%), irrespective of the classification system used. Only 1 patient belonged to the group with an intermediate risk when using the criteria by Fletcher (6.6%) and Joensuu (7.1%). Around 20% belonged to the low and very low risk group and this number was even higher when applying the criteria of Miettinen (28.5%). In consideration of all patients (Table 3), a tumor progression was noted for 41.6–50% of the high risk group, 0–5.6% of the intermediate group, 6.7–7.1% of the low risk group, and 5.9–7.1% of the very low risk group patients, depending on the risk classification that was applied.

Secondary Malignant Lesions in GIST Patients

Various malignancies are known to be associated with an increased rate of malignant tumors at other, unrelated sites. Secondary neoplasias may precede or succeed the index admission for GIST or are detected simultaneously. We studied how many GIST patients suffered from a secondary neoplasm and which kind of tumor they had developed during the same observation period; 44 patients (42.3%) were diagnosed with a secondary tumor of any kind (benign, semi-malignant, or malignant) in addition to the GIST. In 32 patients (30.8%), 38 malignant tumors were found. Among them, 9 tumors were known for at least 1 year before diagnosis of GIST, while 18 tumors were diagnosed within the time period of ± 1 year of index admission for GIST and 9 malignant tumors were diagnosed at least 1 year after the diagnosis of a GIST. In 2 cases, the exact time of diagnosis of the secondary neoplasm could not be retrieved retrospectively. Among them, 26 individuals had 1 additional malignancy and 6 patients 2 more malignant tumors. Moreover, 15 benign tumors were detected in 14 patients (13.5%) and 3 semi-malignant tumors (basal cell carcinoma) were found in 3 patients (2.9%). Combinations of either benign, semi-malignant, or malignant tumors existed in 5 patients.

The number and site of malignant neoplasias are shown in Table 6. Carcinomas of the gastrointestinal tract were observed most frequently (52.6%) followed by tumors of the urogenital tract (21.1%) and the mammary gland (18.4%). Regarding organ involvement, secondary malignancies were most often seen in the colon (26.3%) and the stomach and the prostate (15.8%). Rarer entities were hematologic neoplasias (5.3%) and a peripheral nerve sheath tumor (2.6%), a type of sarcoma originating from the surrounding tissue of nerves.

Table 6. Occurrence and site of secondary malignant neoplasms in our cohort and comparison to literature

		Our cohort number of tumors, <i>n</i> (%)	Pandurengan et al. [1], 2010, <i>n</i> (%)	Agaimy et al. [41], 2006, <i>n</i> (%)
Carcinoma of the urogenital tract	All	8 (21.1)	62 (33)	111 (23)
	Prostate cancer	6 (15.8)	28 (15)	43 (9)
	Kidney cancer	1 (2.6)	12 (6.5)	27 (6)
	Urothelial cancer	1 (2.6)	7 (3.8)	10 (2)
Carcinoma of the gastrointestinal tract	All	20 (52.6)	48 (26)	228 (47)
	Colorectal cancer	10 (26.3)	18 (9.7)	109 (22)
	Pancreatic cancer	3 (7.9)	5 (2.7)	11 (2)
	Gastric cancer	6 (15.8)	5 (2.7)	95 (19)
	Gallbladder cancer	1 (2.6)	1 (0.5)	4 (1)
Breast cancer		7 (18.4)	15 (8)	34 (7)
Hematologic neoplasia	Lymphoma/ leukemia	2 (5.3)	12 (6.5)	36 (7)
Malignant peripheral nerve sheath tumor		1 (2.6)	NA	NA
Total number of additional malignancies		38 (100)	186 (100)	518 (100)

NA, not available.

Discussion

GISTs are common mesenchymal neoplasias of the gastrointestinal tract. Within the last decades, much progress has been made not only in understanding the molecular biology of these tumors but also regarding diagnostic and therapeutic options [25].

Here we characterized GIST patients from a tertiary care hospital in north-eastern Germany diagnosed between 1993 and 2011. On the one hand, we focused on clinical symptoms of these patients in relation to the tumor localization, and on the other hand, we studied metastasis formation and the association with secondary malignancies.

In our cohort, the calculated annual incidence of GISTs was between 0.4 and 2.6 per 100,000 and thus in line with reported data from other groups, especially from north-western Europe (The Netherlands, Sweden, Iceland) where it ranged from 0.7 to 2.0 per 100,000 [7, 26, 27]. Since our hospital is the only tertiary referral center in Western Pomerania, a rural area in north-eastern Germany with a population of around 700,000 inhabitants [23], it is a well suited area to make near population-based assessments, because nearly all patients are referred to this hospital. Secondly, the study population was well defined and histology was available for patients from the same team of pathologists. In our cohort, the mean age at first diagnosis of GIST was slightly higher (66.86 years)

compared to other series reporting a range between 57.8 and 65 years [28–30]. This may be due to a slightly older population in north-eastern Germany compared to the rest of the country being more than 3 years above the mean [31–34]. Another reason for the later age at diagnosis may be a more limited access to specialist care in this rural area.

As outlined by the AFIP tumor size, mitotic index and location are prognostic factors for this tumor entity. In our study, the mean tumor size was 4.96 cm and thus quite comparable to another population-based study on GISTs from Europe [27]. Around two-thirds (67.6%) were located in the stomach, consistent to other groups who indicated gastric GISTs being the most frequent stromal tumor in 40–60% of cases. Tumor manifestation in the small intestine was seen in 16.2%, compared to studies that found jejunal or ileal stromal tumors in 20–50% of patients [15, 19, 35]. In contrast to the small intestine, a colorectal manifestation is rarely observed having an incidence of 4.0–6.4% and thus slightly higher than in our cohort (2.9%) [15, 19, 21, 35]. Esophageal location was comparably very low to other studies (mostly <5%), the same was true for extraintestinal GISTs. Data for extraintestinal GISTs were rather scarce. In a large monocentric study of 200 patients with GIST, an extraintestinal location was seen in 8% whereby they also included esophageal tumors in that entity [29].

A considerable number of our patients (around 30%) remained clinically asymptomatic. In these patients, diagnosis was often established either by radiologic, ultrasound, or endoscopic examinations done for reasons other than a suspected tumor, during an operation, or in surgical specimens after bariatric surgery. The frequency of asymptomatic GIST patients is in line with other studies that reported an incidence of 10–30% [35]. Establishing the diagnosis of GIST is still challenging as these tumors do not have a specific clinical manifestation. Moreover, symptoms depend on tumor size and can be observed in other gastrointestinal disorders as well [36]. In addition, some patients present with more than 1 clinical sign. In our cohort, common symptoms were gastrointestinal bleeding including either hematemesis, hematochezia, melena or a positive stool blood test (55.8%), abdominal pain (38.5%), weight loss, or a general feeling of weakness and dizziness (each 13.5%). These observations were in line with other series that found comparable results [15, 21, 36, 37].

We were further interested in whether specific clinical symptoms of GIST patients may predict tumor location. For gastric GISTs, the largest subgroup, gastrointestinal bleeding was by far the most common reported symptom (64%) and more frequently observed than for other GIST locations. Abdominal pain was less often seen when compared to all GISTs together (28 vs. 38.5%). The absence of clinical symptoms was often noted when tumor site was in the esophagus, small, or large intestine. However, numbers of patients in the latter subgroups were rather small so that these results have to be interpreted with caution. Our results confirm observations from Caterino et al. [21] who found that bleeding in the digestive tract was more frequent in gastric GISTs than for other localizations. Abdominal pain was observed almost as often as bleeding which differs to our results [21]. However, their patient cohort was smaller. In addition, as in our study, the authors were also confronted with a low case load for extra-gastric stromal tumors making representative statistics more difficult.

An accurate prediction of the biological behavior of GISTs is challenging. The majority of our tumors were classified with a low or very low recurrence risk independent of the type of classification that was used. Around one-quarter of patients belonged to the high risk group (24.1–27.6%). Regarding metastasis formation only 15 out of 104 (14.4%) patients were diagnosed with metastatic disease. Among them two-thirds had metachronous and one-third synchronous metastases. We compared the presence of metastases to the most commonly used risk classification systems in order to see accuracy of prediction. As expected, more than 70% of patients with metastases belonged to the

high risk group. Only a minority of patients were classified as intermediate or very low risk. In contrast to our expectations, our data show that even 13.3–21.4% of metastasized patients belonged to the low risk group.

A recently published large single center study ($n = 497$) investigating tumor recurrence in GIST patients found a slightly higher risk of overall tumor recurrence or metastasis formation which was 19.7% [38]. However, hardly any tumor relapse was observed in the low risk and very low risk groups. There may be various reasons explaining this discrepancy. One is that our study contained a higher proportion of low and very low risk GIST cases. Secondly, there may be differences regarding the time of tumor diagnosis, operation, and adjuvant chemotherapy between the centers that may influence recurrence rates. Further studies will be necessary to elucidate this question. In addition, the absolute number of patients with metastasis formation or tumor recurrence was low in the low risk categories making a universal conclusion difficult. In a large meta-analysis of population-based studies including 2,560 GIST patients and a validation series of 920 patients, Joensuu et al. [39] identified the modified NIH classification as most suitable criteria for identification of high risk groups among the commonly used classification systems. Secondly, they developed a new method for estimation of GIST recurrence by including tumor localization and rupture status and respecting the non-linear effect of mitosis and tumor size on recurrence-free survival. Our data showed that NIH criteria predicted tumor recurrence (73.3%) slightly better than modified NIH criteria and AFIP criteria (71.4%); however, only 15 of our patients had metastases.

Besides metastasis formation, the occurrence of secondary neoplasms worsens the prognosis of GIST patients. There is evidence that coexisting secondary neoplasms appear more often in GIST patients than in the general population [1, 40] and an association of GISTs with both solid and hematologic tumors were reported. The overall frequency of secondary tumors varies from 4.5 to 33% with a mean of 13% [1, 41]. According to Agaimy et al. [41], gastric cancer (47%), prostate cancer (9%), breast cancer or lymphoma, as well as leukemia (both 7%) were reported in association with GISTs, followed by kidney (6%) and lung cancer (5%) as well as tumors of the female reproductive tract (5%). In a large single center study of 783 patients, 20% were diagnosed with at least 1 additional primary [1] and the majority of tumors were of genitourinary (33%) or gastrointestinal (26%) origin. To a lesser extent, breast cancer (8%) and hematologic neoplasias (6.5%) were seen. In our series, more than 40% of GIST patients were diagnosed with an

additional tumor, irrespective of dignity. A malignant secondary neoplasia was observed in 30.8% and thus being in the upper range of reported prevalences. Three patients (2.9%) had diagnosis of both a benign and a malignant neoplasia. Potential reasons for our comparatively high number of secondary neoplasms are a longer follow-up time in our study that was up to 15 years and a much higher mean age of our study population which was around 67 years at the first diagnosis of GIST, and thus more than 10 years older than in the aforementioned study [1]. In addition, our patient group was exclusively of Caucasian ancestry. Our data support the high incidence of secondary GI neoplasms in GIST patients, especially colorectal, gastric, and pancreatic cancer.

There are limitations of our study. The retrospective setting caused missing data that we had to correct for when analyzing histopathology and clinical symptoms. Information on clinical signs was unavailable for about 25% of patients. Therefore, a subgroup analysis of symptoms depending on GIST localization is challenging due to small numbers. In addition, reporting of patient's complaints in a retrospective analysis has limitations as they have to be retrieved from the patient file which might be incomplete. Reporting of symptoms in a prospective design would be a superior approach. Secondly, a comprehensive characterization of GISTs needs to include an analysis of genetic changes that are likely to further affect survival [42]. Larger, preferably multicenter studies will be necessary to determine symptoms depending on tumor location, especially for extra-gastric tumor sites. Moreover, the small number of patients limits the calculation of probability of disease recurrence after surgery, since follow-up data were available for only 39 patients and ex-

act time of death was known for 12 patients. Metachronous metastases were detected in only 10 patients. Larger studies with pooled data will be an ideal approach for estimation of GIST recurrence as published previously [39].

In summary, we presented a clinical characterization of a single center cohort of GIST patients. Clinical symptoms correlated with tumor location, most remarkably for gastric location. There was a considerable risk for tumor recurrence, metastasis formation, and the occurrence of secondary malignant tumors. A comparably higher risk for metastases was detected especially in low risk patients. These observations have clinical implications. First, even patients in lower risk groups may benefit from follow-up examinations, although it remains unclear whether this is cost-effective, and secondly, GIST patients with their increased risk of developing secondary tumors at unrelated sites may benefit from surveillance and screening programs for colorectal, breast, and prostate cancer.

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Disclosure Statement

The authors declare no conflicts of interest.

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